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#### Amendments to the Claims

# 1. (currently amended) A compound of formula I

$$\begin{array}{c|cccc}
R^8 & R^1 & N \\
R^9 & R^{10} & R^5 \\
R & A & I
\end{array}$$

or a pharmaceutically acceptable salt, crystal-form, or hydrate, wherein:

### A is

- a) an aryl ring <u>selected from phenyl</u>, wherein any stable <u>phenyl</u> aryl ring atom is independently unsubstituted or substituted with
  - 1) halogen,
  - 2) NO<sub>2</sub>,
  - 3) CN,
  - 4)  $CR^{46}=C(R^{47}R^{48})_2$ ,
  - 5)  $C \equiv C R^{46}$ ,
  - 6) (CRiRJ)rOR46
  - 7)  $(CR^{i}R^{j})_{r}N(R^{46}R^{47})$ ,
  - 8)  $(CR^iR^j)_r C(O)R^{46}$ ,
  - 9)  $(CR^iR^j)_r C(O)OR^{46}$ ,
  - 10)  $(CR^{i}R^{j})_{r}R^{46}$ ,
  - 11)  $(CR^{i}R^{j})_{r} S(O)_{0-2}R^{61}$ ,
  - 12)  $(CR^{i}R^{j})_{r} S(O)_{0-2}N(R^{46}R^{47}),$
  - 13)  $OS(O)_{0-2}R^{61}$ ,
  - 14) N(R<sup>46</sup>)C(O)R<sup>47</sup>,
  - 15) N(R46)S(O)0-2R61,
  - 16)  $(CR^{i}R^{j})_{r}N(R^{46})R^{61}$ ,
  - 17) (CRiRJ)<sub>r</sub>N(R<sup>46</sup>)R<sup>61</sup>OR<sup>47</sup>,
  - 18)  $(CR^{i}R^{j})_{r}N(R^{46})(CR^{k}R^{l})_{s}C(O)N(R^{47}R^{48}),$
  - 19) N(R46)(CRiRj)<sub>r</sub>R61,
  - 20)  $N(R^{46})(CR^{i}R^{j})_{r}N(R^{47}R^{48})$ ,
  - 21)  $(CR_iR_i)_rC(O)N(R_i^{47}R_i^{48})$ ,



22) oxo,

b) a heteroaryl ring <u>selected from the group consisting of pyridine</u>, <u>pyrimidine</u>, <u>pyrimidine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyridine</u>, <u>pyrimidine</u>, <u>pyri</u>

a 5-membered unsaturated monocyclic ring with 1, 2, 3 or 4 heteroatom ring atoms selected from the group consisting or N, O or S, a 6-membered unsaturated monocyclic ring with 1, 2, 3 or 4 heteroatom ring atoms selected from the group consisting N, O and S, and

a 9-or 10-membered unsaturated bicyclic ring with 1, 2, 3 or 4 heteroatom ring atoms selected from the group consisting or N, O or S,

wherein any stable S heteroaryl ring atom is unsubstituted or mono- or di-substituted with oxo, and any stable C or N heteroaryl ring atom is independently unsubstituted or substituted with

- 1) halogen,
- 2) NO<sub>2</sub>,
- 3) CN,
- 4)  $CR^{46}=C(R^{47}R^{48})_2$ ,
- 5)  $C = C R^{46}$ ,
- 6) (CRiRI)<sub>r</sub>OR<sup>46</sup>,
- 7)  $(CR^{i}R^{j})_{r}N(R^{46}R^{47})$ ,
- 8)  $(CR^{i}R^{j})_{r} C(O)R^{46}$ ,
- 9)  $(CR^iR^j)_r C(O)OR^{46}$ ,
- 10) (CRiRJ)<sub>r</sub>R<sup>46</sup>,
- 11)  $(CR^{i}R^{j})_{r} S(O)_{0-2}R^{61}$ ,
- 12)  $(CR^{i}R^{j})_{r} S(O)_{0-2}N(R^{46}R^{47}),$
- 13)  $OS(O)_{0-2}R^{61}$ ,
- 14) N(R<sup>46</sup>)C(O)R<sup>47</sup>,
- 15)  $N(R^{46})S(O)_XR^{61}$ ,
- 16)  $(CR^{i}R^{j})_{r}N(R^{46})R^{61}$ ,
- 17)  $(CR^{i}R^{j})_{r}N(R^{46})R^{61}OR^{47}$ ,
- 18)  $(CR^{i}R^{j})_{r}N(R^{46})(CR^{k}R^{l})_{s}C(O)N(R^{47}R^{48}),$
- 19) N(R<sup>46</sup>)(CR<sup>i</sup>R<sup>j</sup>)<sub>r</sub>R<sup>61</sup>,
- 20)  $N(R^{46})(CR^{i}R^{j})_{r}N(R^{47}R^{48})$ ,
- 21)  $(CR^{i}R^{j})_{r}C(O)N(R^{47}R^{48})$ , or



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## 22) oxo, or

c) a 4-, 5- or 6-membered heterocyclic ring containing 1 or 2 nitrogen atoms, unsubstituted, mono-substituted or di-substituted with C<sub>1</sub>-C<sub>6</sub> alkyl;

Y is CH<sub>2</sub>, NR<sup>53</sup>, NC(O)R<sup>53</sup>, S(O)<sub>0-2</sub> or O;

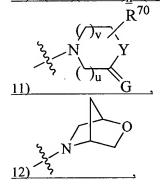
G is H<sub>2</sub> or O;

Ra, Rb, Re, Rd, Re, Rf, Rg, Rh, Ri, Rj, Rk, and Rl are independently selected from the group consisting of:

- 1) hydrogen,
- 2) C1-C6 alkyl,
- 3) halogen,
- 4) aryl,
- 5)  $R^{80}$ ,
- 6) C3-C10 cycloalkyl, and
- 7) OR4,

said alkyl, aryl, and cycloalkyl being unsubstituted, monosubstituted with  $R^7$ , disubstituted with  $R^7$  and  $R^{15}$ , trisubstituted with  $R^7$ ,  $R^{15}$  and  $R^{16}$ , or tetrasubstituted with  $R^7$ ,  $R^{15}$ ,  $R^{16}$  and  $R^{17}$ ;  $R^1$  is independently selected from:

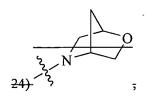
- 1) hydrogen,
- 2) halogen,
- 3) CN,
- 4) OR40
- 5)  $N(R^{40}R^{41})$
- 6)  $C(O)OR^{40}$ ,
- 7) R81,
- 8) S(O)0-2R6,
- 9)  $N(R^{40})(CR^{a}R^{b})_{n}R^{6}$ , wherein  $R^{6} = R^{83}$ ,
- 10) N(R<sup>40</sup>)(CRaRb)<sub>n</sub>N(R<sup>41</sup>R<sup>42</sup>),



# 13) C(O)N(R41R42), and

14) a 4-, 5-, or 6-membered heterocyclic ring containing 1 nitrogen atom, unsubstituted, or mono-, di- or tri-substituted with -OH-

- 1) hydrogen,
- 2) halogen,
- 3) NO<sub>2</sub>,
- 4<del>) CN,</del>
- 5)-CR40-C(R41R42),
- 6)  $C = CR^{40}$
- 7) (CRaRb)<sub>n</sub>OR<sup>40</sup>.
- 8) (CRaRb) nN(R40R41),
- 9)  $(CR^aR^b)_nC(O)R^{40}$ ,
- 10) (CRaRb)<sub>n</sub>C(O)OR40,
- 11) (CRaRb)nR40;
- $12) (CR^aR^b)_nS(O)_{0-2}R^6$
- $13) (CR^aR^b)_nS(O)_{0-2}N(R^{40}R^{41}),$
- 14) OS(O)0-2R6,
- $15) N(R^{40})C(O)R^{41}$
- $16) N(R^{40})S(O)_{0-2}R^{6}$
- 17) (CRaRb)nN(R40)R6,
- 18) (CRaRb)nN(R40)R6OR41,
- 19) (CRaRb)nN(R40)(CReRd)tC(O)N(R41R42),
- 20) N(R40)(CRaRb)nR6,
- $21) N(R^{40})(CR^aR^b)_n N(R^{41}R^{42})_7$



25) (CRaRb)nC(O)N(R41R42), and

26) a 4-, 5-, or 6-membered heterocyclic ring containing 1 nitrogen atom, unsubstituted, or mono-, di- or tri-substituted with OH;

R<sup>2</sup>, R<sup>8</sup>, R<sup>9</sup>-and R<sup>10</sup> are independently selected from <u>hydrogen and halogen</u>; R<sup>9</sup> is OCH<sub>3</sub> or OCHF<sub>2</sub>.

- 1)-hydrogen,
- 2) halogen,
- 3) NO2;
- 4) CN,
- 5) CR43=C(R44R45),
- 6) C=CR43,
- 7) (CReRf) pOR43
- 8) (CReRf)pN(R43R44),
- 9)  $(CReRf)_pC(O)R43$ ,
- 10) (CReRf)<sub>p</sub>C(O)OR43,
- 11) (CReRf)<sub>D</sub>R43,
- 12) (CReRf)<sub>p</sub>S(O)<sub>0-2</sub>R60,
- 13) (CReRf)<sub>p</sub>S(O)<sub>0-2</sub>N(R43R44),
- $14) OS(O)_{0-2}R^{60}$
- 15) N(R43)C(O)R44,
- $16) N(R^{43})S(O)_{0-2}R^{60}$
- 17) (CReRf)<sub>D</sub>N(R43)R60,
- 18) (CReRf)<sub>p</sub>N(R43)R60OR44,
- $49) (CR^{e}R^{f})_{p}N(R^{43})(CR^{g}R^{h})_{q}C(O)N(R^{44}R^{45}),$
- $20) N(R^{43})(CReR^{f})_{p}R^{60}$
- 21) N(R43)(CReRf), N(R44R45), and
- 22) (CReRf)<sub>p</sub>C(O)N(R43R44),
- or R<sup>2</sup> and R<sup>8</sup> are independently as defined above, and R<sup>9</sup> and R<sup>10</sup>, together with the atoms to which they are attached, form the ring



R4, R40, R41, R42, R43, R44, R45, R46, R47, R48, R49, R50, R51, R52, and R53 are independently selected from:

- 1) hydrogen,
- 2) C<sub>1</sub>-C<sub>6</sub> alkyl,
- 3) C3-C10 cycloalkyl,
- 4) aryl,
- 5) R81,
- 6) CF<sub>3</sub>,
- 7) C2-C6 alkenyl, and
- 8) C2-C6 alkynyl,

said alkyl, aryl, and cycloalkyl is unsubstituted, mono-substituted with R<sup>18</sup>, di-substituted with R<sup>18</sup> and R<sup>19</sup>, tri-substituted with R<sup>18</sup>, R<sup>19</sup> and R<sup>20</sup>, or tetra-substituted with R<sup>18</sup>, R<sup>19</sup>, R<sup>20</sup> and R<sup>21</sup>; R<sup>5</sup> is independently selected from:

- 1) hydrogen,
- 2) halogen,
- 3) CN,
- 4)  $C(O)N(R^{49}R^{50})$ ,
- 5)  $C(O)OR^{49}$ ,
- 6)  $S(O)_{0-2}N(R^{49}R^{50})$ ,
- 7)  $S(O)_{0-2}R^{62}$ ,
- 8) C1-C6 alkyl,
- 9) C3-C10 cycloalkyl,
- 10) R82,

said alkyl, aryl, and cycloalkyl is unsubstituted, mono-substituted with  $R^{22}$ , di-substituted with  $R^{22}$  and  $R^{23}$ , tri-substituted with  $R^{22}$ ,  $R^{23}$  and  $R^{24}$ , or tetra-substituted with  $R^{22}$ ,  $R^{23}$ ,  $R^{24}$  and  $R^{25}$ ;  $R^{60}$ ,  $R^{61}$ ,  $R^{62}$  and  $R^{63}$  are independently selected from:

- 1) C1-C6 alkyl,
- 2) aryl,
- 3) R83, and
- 4) C3-C10 cycloalkyl;

said alkyl, aryl, and cycloalkyl is unsubstituted, mono-substituted with  $R^{26}$ , di-substituted with  $R^{26}$  and  $R^{27}$ , tri-substituted with  $R^{26}$ ,  $R^{27}$  and  $R^{28}$ , or tetra-substituted with  $R^{26}$ ,  $R^{27}$ ,  $R^{28}$  and  $R^{29}$ ;



R7, R15, R16, R17, R18, R19, R20, R21, R22, R23, R24, R25, R26, R27, R28, R29, and R70 are independently selected from:

- 1) C<sub>1</sub>-C<sub>6</sub> alkyl,
- 2) halogen,
- 3) OR51,
- 4) CF<sub>3</sub>,
- 5) aryl,
- 6) C3-C10 cycloalkyl,
- 7) R84,
- 8)  $S(O)_{0-2}N(R^{51}R^{52})$ ,
- 9) C(O)OR<sup>51</sup>,
- 10)  $C(O)R^{51}$ ,
- 11) CN,
- 12) C(O)N(R<sup>5</sup>1R<sup>5</sup>2),
- 13)  $N(R^{51})C(O)R^{52}$ ,
- 14)  $S(O)_{0-2}R^{63}$ ,
- 15) NO2, and
- 16) N(R<sup>51</sup>R<sup>52</sup>);

R80, R81, R82, R83 and R84 are independently selected from a group of unsubstituted or substituted heterocyclic rings consisting of a 4-6 membered unsaturated or saturated monocyclic ring with 1, 2, 3 or 4 heteroatom ring atoms selected from the group consisting N, O and S, and a 9- or 10-membered unsaturated or saturated bicyclic ring with 1, 2, 3 or 4 heteroatom ring atoms selected from the group consisting or N, O or S;

n,  $\frac{1}{p}$ ,  $\frac{1}{q}$ , r, s and t are independently 0, 1, 2, 3, 4, 5 or 6; u is 0, 1 or 2; and v is 0, 1 or 2.

- 2.(canceled).
- 3. (canceled).
- 4. (original) A compound of Claim <u>1</u>-3, or a pharmaceutically acceptable salt thereof, wherein R<sup>1</sup> is selected from the group consisting of hydrogen, -SCH<sub>3</sub>, -SO<sub>2</sub>CH<sub>3</sub>, -NH(CH<sub>2</sub>)<sub>3</sub>OH, -NH(CH<sub>2</sub>)<sub>2</sub>OCH<sub>3</sub>,



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-NH(CH<sub>2</sub>)<sub>3</sub>OCH<sub>3</sub>, -NH(CH<sub>2</sub>)<sub>2</sub>NH<sub>2</sub>, -NH<sub>2</sub>, -SO<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>, -CN, Cl, -OCH<sub>3</sub>,

- -OCH<sub>2</sub>CHCH<sub>2</sub>, -OCH<sub>2</sub>CH(OH)CH<sub>2</sub>OH, -NHCH<sub>2</sub>CHCH<sub>2</sub>, -CH<sub>3</sub>, -CH<sub>2</sub>CH<sub>2</sub>OH,
- $-O(CH_2)_2CHCH_2, -O(CH_2)_2CH(OH)(CH_2OH), -NHCH(CH_2OH)_2, \\$
- -NHCH<sub>2</sub>CH(OH)CH<sub>2</sub>OH, -NH(CH<sub>2</sub>)<sub>2</sub>CH(OH)CH<sub>2</sub>OH,

$$-NHCH_{2} \longrightarrow OCH_{3}$$

$$-OCH_{2} \longrightarrow OCH_{2} \longrightarrow O$$

5. (original) A compound of Claim 4, or a pharmaceutically acceptable salt thereof, wherein A is selected from the group consisting of

- 1) phenyl, wherein any stable ring atom is unsubstituted or substituted with halogen,
- 2) pyridinyl, wherein any stable C ring atom is unsubstituted or substituted with halogen,
- 3) indolyl, wherein any stable C or N ring atom is unsubstituted or substituted with halogen, and
- 4) a heterocyclic ring selected from the group consisting of pyrrolidine, piperidine, piperazine, and azetidine, unsubstituted, mono-substituted or di-substituted with C<sub>1</sub>-C<sub>6</sub> alkyl.

6. (original) A compound of Claim 5, or a pharmaceutically acceptable salt thereof, wherein R<sup>5</sup> is selected from the group consisting of CN and C<sub>1</sub>-C<sub>6</sub> alkyl, wherein said alkyl is unsubstituted, mono-substituted with R<sup>22</sup>, di-substituted with R<sup>22</sup> and R<sup>23</sup>, tri-substituted with R<sup>22</sup>, R<sup>23</sup> and R<sup>24</sup>, or tetra-substituted with R<sup>22</sup>, R<sup>23</sup>, R<sup>24</sup> and R<sup>25</sup>.

7. (original) A compound of Claim 6, or a pharmaceutically acceptable salt thereof, selected from the group consisting of [(6-methoxy-4-phenylisoquinolin-3-yl)methyl]dimethylamine,

1-(1-chloro-6-methoxy-4-phenylisoquinolin-3-yl)-N,N-dimethylmethanamine,



{[6-methoxy-1-(methylthio)-4-phenylisoquinolin-3-yl]methyl}dimethylamine,

[6-methoxy-1-(methylsulfonyl)-4-phenylisoquinolin-3-yl]methyl(dimethyl)amine oxide,

1-[6-methoxy-1-(methylsulfonyl)-4-phenylisoquinolin-3-yl]-N,N-dimethylmethanamine,

3-[(dimethylamino)methyl]-6-methoxy-4-phenylisoquinoline-1-carbonitrile,

2,3-Dimethyl-6-methoxy-4-phenylisoquinolinium hydroxide,

6-methoxy-1-(2-methoxyethoxy)-3-methyl-4-phenylisoquinoline,

{3-[(6-methoxy-3-methyl-4-phenylisoquinolin-1-yl)oxy]propyl}amine,

2-[(6-methoxy-3-methyl-4-phenylisoquinolin-1-yl)amino]ethanol,

6-methoxy-3-methyl-1-(methylsulfonyl)-4-phenylisoquinoline,

6-methoxy-N-(2-methoxyethyl)-3-methyl-4-phenylisoquinolin-1-amine,

N-(6-methoxy-3-methyl-4-phenylisoquinolin-1-yl)ethane-1,2-diamine,

6-methoxy-3-methyl-4-phenylisoquinoline,

N-(3,4-dimethoxybenzyl)-6-methoxy-3-methyl-4-phenylisoquinolin-1-amine,

6-methoxy-3-methyl-4-phenylisoquinolin-1-amine,

1-(ethylsulfonyl)-6-methoxy-3-methyl-4-phenylisoquinoline,

1-(benzylsulfonyl)-6-methoxy-3-methyl-4-phenylisoquinoline,

6-methoxy-3-methyl-4-phenyl-1-(phenylsulfonyl)isoquinoline,

6-methoxy-3-methyl-4-phenylisoquinoline-1-carbonitrile,

3-tert-butyl-6-methoxy-1-(2-methoxyethoxy)-4-phenylisoquinoline,

1-chloro-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

6-methoxy-4-phenylisoquinoline-1,3-dicarbonitrile,

1-(allyloxy)-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-(2,3-dihydroxypropoxy)-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(allylamino)-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(+/-)-1-[(2,3-dihydroxypropyl)amino]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-{[(2S)-2,3-dihydroxypropyl]amino}-6-methoxy-4-phenylisoquinoline-3-carbonitrile,



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1-{[(2R)-2,3-dihydroxypropyl]amino}-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(+/-)-1-[(2,2-dimethyl-1,3-dioxolan-4-yl)methoxy]-6-methoxy-4-phenylisoquinoline-3-

carbonitrile,

 $1-\{[(4S)-2,2-dimethyl-1,3-dioxolan-4-yl]methoxy\}-6-methoxy-4-phenylisoquinoline-3-methoxy-4$ 

carbonitrile,

1-{[(4R)-2,2-dimethyl-1,3-dioxolan-4-yl]methoxy}-6-methoxy-4-phenylisoquinoline-3-

carbonitrile,

1-{[(2R)-2,3-dihydroxypropyl]oxy}-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-{[(2S)-2,3-dihydroxypropyl]oxy}-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(+/-)-1-{[2,3-dihydroxypropyl]oxy}-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-[(3R)-3-hydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-[(3S)-3-hydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(+/-)-1-[3-hydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-[cis-3,4-dihydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

6-methoxy-4-phenyl-1-pyrrolidin-1-ylisoquinoline-3-carbonitrile,

6-methoxy-1-(methylsulfonyl)-4-phenylisoquinoline-3-carbonitrile,

6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1,6-dimethoxy-4-phenylisoquinoline-3-carbonitrile,

1-chloro-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

4-(3-fluorophenyl)-6-methoxy-1-methylisoquinoline-3-carbonitrile,

4-(3-fluorophenyl)-1-[(2-hydroxyethyl)amino]-6-methoxyisoquinoline-3-carbonitrile,

1-amino-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

4-(3-fluorophenyl)-1-[(3-hydroxypropyl)amino]-6-methoxyisoquinoline-3-carbonitrile,

1-(but-3-enyloxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

(+/-)-1-(2,3-dihydroxypropoxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,



1-[(2R)-2,3-dihydroxypropoxy]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

1-[(2S)-2,3-dihydroxypropoxy]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

(+/-)-1-(3,4-dihydroxybutoxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

(+/-)-1-[(3R)-3,4-dihydroxybutoxy]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

1-[(3S)-3,4-dihydroxybutoxy]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

(+/-)-1-[(1,4-dioxan-2-ylmethyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(1,4-dioxan-(2R)-2-ylmethyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(1,4-dioxan-(2S)-2-ylmethyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

4-(3-fluorophenyl)-6-methoxy-1-[(1-methyl-1H-imidazol-4-yl)methoxy]isoquinoline-3-

carbonitrile,

(+/-)-1-(1,3-dioxolan-4-ylmethoxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-(1,3-dioxolan-(4R)-4-ylmethoxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

1-(1,3-dioxolan-(4S)-4-ylmethoxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

1-(1,3-dioxan-5-yloxy)-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

4-(3-fluorophenyl)-1-{[2-hydroxy-1-(hydroxymethyl)ethyl]amino}-6-methoxyisoquinoline-

3-carbonitrile,

4-(3-fluorophenyl)-1-(1H-imidazol-5-ylmethoxy)-6-methoxyisoquinoline-3-carbonitrile,

1-{[(2R)-2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-{[(2S)-2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

(+/-)-1-{[2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-



1-(1H-imidazol-1-yl)-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

6-methoxy-4-phenyl-1-[(pyridin-2-ylmethyl)amino]isoquinoline-3-carbonitrile,

6-methoxy-4-phenyl-1-[(2-pyridin-2-ylethyl)amino]isoquinoline-3-carbonitrile,

(+/-)-1-[(3,4-dihydroxybutyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(3R)-(3,4-dihydroxybutyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(3S)-(3,4-dihydroxybutyl)amino]-4-(3-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-chloro-4-(2-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

4-(2-fluorophenyl)-6-methoxyisoquinoline-3-carbonitrile,

(+/-)-1-[(2,3-dihydroxypropyl)amino]-4-(2-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(2S)-(2,3-dihydroxypropyl)amino]-4-(2-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

1-[(2R)-(2,3-dihydroxypropyl)amino]-4-(2-fluorophenyl)-6-methoxyisoquinoline-3-

carbonitrile,

(+/-)-6-(difluoromethoxy)-1-{[2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,

6-(difluoromethoxy)-1-{[(2S)-2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,

6-(difluoromethoxy)-1-{[(2R)-2,3-dihydroxypropyl]amino}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,

(+/-)-6-(difluoromethoxy)-1-{[2,3-dihydroxypropyl]oxy}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,



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6-(difluoromethoxy)-1-{[(2S)-2,3-dihydroxypropyl]oxy}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,

6-(difluoromethoxy)-1-{[(2R)-2,3-dihydroxypropyl]oxy}-4-(3-fluorophenyl)isoquinoline-

3-carbonitrile,

1-(4-hydroxypiperidin-1-yl)-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-azetidin-1-yl-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

(+/-)-1-[trans-3,4-dihydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-[(3R,4R)-3,4-dihydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile,

1-[(3S,4S)-3,4-dihydroxypyrrolidin-1-yl]-6-methoxy-4-phenylisoquinoline-3-carbonitrile, and

6-methoxy-N-(3-methoxypropyl)-3-methyl-4-phenylisoquinolin-1-amine.

8. (withdrawn) A method of treating a condition in a mammal, the treatment of which is effected or facilitated by  $K_V 1.5$  inhibition, which comprises administering a compound of Claim 1 in an amount that is effective at inhibiting  $K_V 1.5$ .

- 9. (withdrawn) A method of Claim 8, wherein the condition is cardiac arrythmia.
- 10. (withdrawn) A method of Claim 9, wherein the cardiac arrythmia is atrial fibrillation.
- 11. (withdrawn) A method of Claim 9, wherein the cardiac arrythmia is selected from the group consisting of atrial flutter, atrial arrhythmia and supraventricular tachycardia.
- 12. (withdrawn) A method of preventing a condition in a mammal, the prevention of which is effected or facilitated by  $K_V 1.5$  inhibition, which comprises administering a compound of Claim 1 in an amount that is effective at inhibiting  $K_V 1.5$ .
  - 13. (withdrawn) A method of Claim 12, wherein the condition is cardiac arrythmia.
- 14. (withdrawn) A method of Claim 13, wherein the cardiac arrythmia is atrial fibrillation.



15. (withdrawn) A method of Claim 13, wherein the cardiac arrythmia is selected from the group consisting of atrial flutter, atrial arrhythmia and supraventricular tachycardia.

- 16. (withdrawn) A method of Claim 12, wherein the condition is a thromboembolic event.
- 17. (withdrawn) A method of Claim 16, wherein the thromboembolic event is a stroke.
- 18. (withdrawn) A method of Claim 12, wherein the condition is congestive heart failure.
- 19. (currently amended) A pharmaceutical formulation comprising a pharmaceutically acceptable carrier and the compound Claim 1 or a pharmaceutically acceptable salt erystal form or hydrate thereof.
- 20. (original) A pharmaceutical composition made by combining the compound of Claim 1 and a pharmaceutically acceptable carrier.
- 21. (withdrawn) A method of treating cardiac arrythmia comprising administering a compound of Claim 1 with a compound selected from one of the classes of compounds consisting of antiarrhythmic agents having Kv1.5 blocking activities, ACE inhibitors, angiotensin II antagonists, cardiac glycosides, L-type calcium channel blockers, T-type calcium channel blockers, selective and nonselective beta blockers, endothelin antagonists, thrombin inhibitors, aspirin, nonselective NSAIDs, warfarin, factor Xa inhibitors, low molecular weight heparin, unfractionated heparin, clopidogrel, ticlopidine, IIb/IIIa receptor antagonists, 5HT receptor antagonists, integrin receptor antagonists, thromboxane receptor antagonists, TAFI inhibitors and P2T receptor antagonists.
- 22. (withdrawn) A method for inducing a condition of normal sinus rhythm in a patient having atrial fibrillation, which comprises treating the patient with a compound of Claim 1.
- 23. (withdrawn) A method for treating tachycardia in a patient which comprises treating the patient with an antitachycardia device in combination with a compound of Claim 1.

